



# State Medical Board of Ohio Report of RU-486 Event

(Required pursuant to R.C. 2919.123)

To be completed by the physician who provided RU-486

1. Date RU-486 was provided:	1	29	18
	Month	Day	Year
2. Name of medical practice or facility at which RU-486 was provided: <u>Planned Parenthood of Greater Ohio</u>			
3. Address of medical practice or facility at which RU-486 was provided: <u>25350 Rockside Rd. Bedford Heights, Ohio 44146</u>			
4. Date post RU-486 complication began: <u>2/1/18</u>			
5. Event(s) (Please check all that apply):			
<input checked="" type="checkbox"/> Incomplete abortion <input type="checkbox"/> Adverse reaction to RU-486 <input type="checkbox"/> Patient hospitalized <input type="checkbox"/> Patient received a transfusion <input type="checkbox"/> Severe bleeding <input type="checkbox"/> Other serious event (specify) _____			
6. Duration of event: <u>1</u> Hours <u>    </u> Days			
7. Remarks: <u>Med ab procedure initiated per FDA regimen on 1/29/18. Pt. 40 no results from medication - was given 2nd dose of misoprostol on 2/1/18. Flw ultrasound on 2/6/18 showed failed procedure. Surgical aspiration was done on 2/6/18; pt did well post op.</u>			
8. a. Name of physician who provided RU-486: <u>Timothy Kress, MD</u>			
8. b. Physician's signature: <u>Timothy S Kress MD/DO</u>			
Date: <u>3/16/18</u>			

Send completed forms to: State Medical Board of Ohio

Legal Department  
30 E. Broad St., 3<sup>rd</sup> Floor  
Columbus, OH 43215-6127

MEDICAL BOARD